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Respectfully submitted
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RBM/cb

MARKED UP CLAIMS

25. A stable pharmaceutical composition comprising a mixture of

(i) an ibuprofen medicament;

(ii) a domperidone medicament; and

(iii) a carrier material

characterised in that the carrier material is substantially free of povidone [10]and comprises at least one diluent combined with at least one [release modifying agent] disintegrating agent, excluding

(a) a compressed tablet comprising granulated ibuprofen and a carrier - material consisting essentially of either maize starch at 35-38% total tablet weight in combination with dried maize starch at 3-4% total tablet weight or microcrystalline cellulose at 10-11% total tablet weight in combination with croscarmellose sodium at 14-16% total tablet weight and pre-gelled starch at 10% total tablet weight;

(b) a direct compression tablet comprising a carrier material consisting essentially of microcrystalline cellulose at 8-11% total tablet weight and lactose at 5-6% total tablet weight;

(c) a hard gelatin capsule comprising a carrier consisting essentially of maize starch at 15-20% total capsule contents weight in combination with pre-gelled starch at 5-6% total capsule contents weight.

28. A composition according to claim [27] 38 characterised in that the granulating agent is hydroxypropyl cellulose or hydroxypropyl methylcellulose.

33. A composition according to claim 25 in which the ibuprofen medicament is racemic ibuprofen or S(+)-ibuprofen or the [8] sodium or lysine salts thereof, present to an extent of 50-65% by weight of the composition and the domperidone medicament is domperidone or the maleate salt thereof, present to an extent of 1-5% of the composition.